

CLAIMS

1. A microemulsion pre-concentrate comprising a difficultly soluble active agent and a carrier medium comprising
 - 1) a hydrophilic phase which comprises dimethylisosorbide and/or a lower alkyl alkanoic ester,
 - 2) a lipophilic phase, and
 - 3) a surfactant.
2. A composition of claim 1 wherein the active agent is a cyclosporin or a macrolide.
3. A composition as claimed in claim 1 or claim 2, wherein the active agent is selected from Cyclosporin A, rapamycin, 40-O-(2-hydroxy)ethyl rapamycin, 33-epi-chloro-33-desoxy-ascomycin, FK 506 or ascomycin.
4. A composition as claimed in any preceding claim wherein the hydrophilic phase comprises ethyl acetate as lower alkyl alkanoic ester.
5. A composition as claimed in one of claims 1 to 4 for oral or parenteral administration.
6. A pharmaceutical composition for enteral or parenteral administration comprising a macrolide and an acid.
7. A composition as claimed in claim 6 wherein the acid is a mono-, di- or tri-carboxylic acid.
8. A composition as claimed in claim 6 or claim 7 wherein the acid is selected from malonic acid, oxalic acid, citric acid and lactic acid.
9. Use of an acid to stabilise a macrolide in a pharmaceutical composition.
10. A method of stabilising a macrolide in a pharmaceutical composition, which method comprises mixing an acid with the macrolide.

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